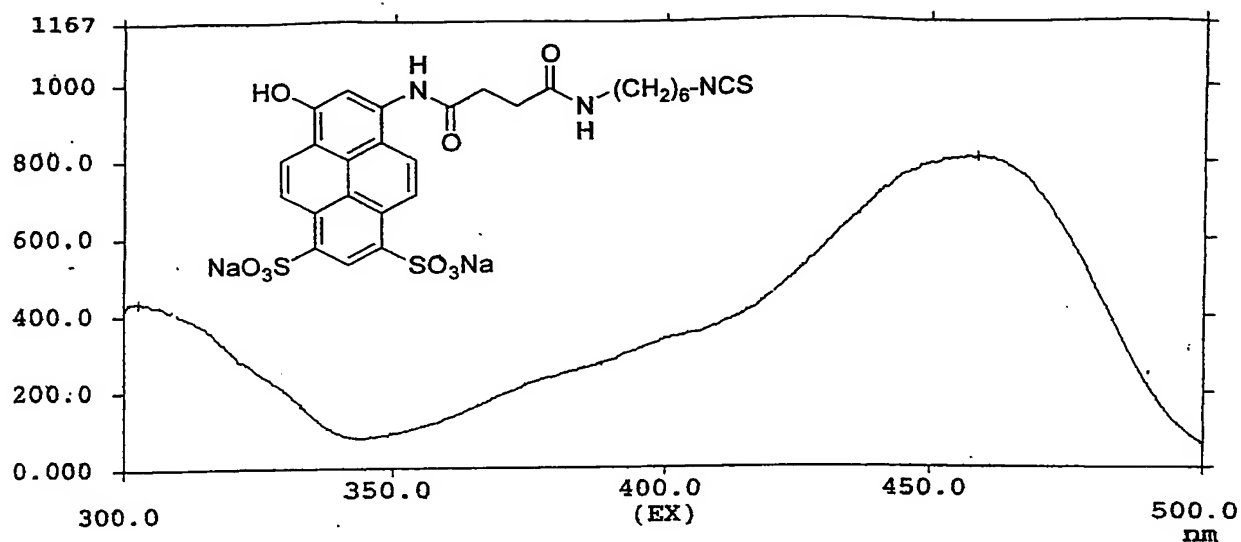


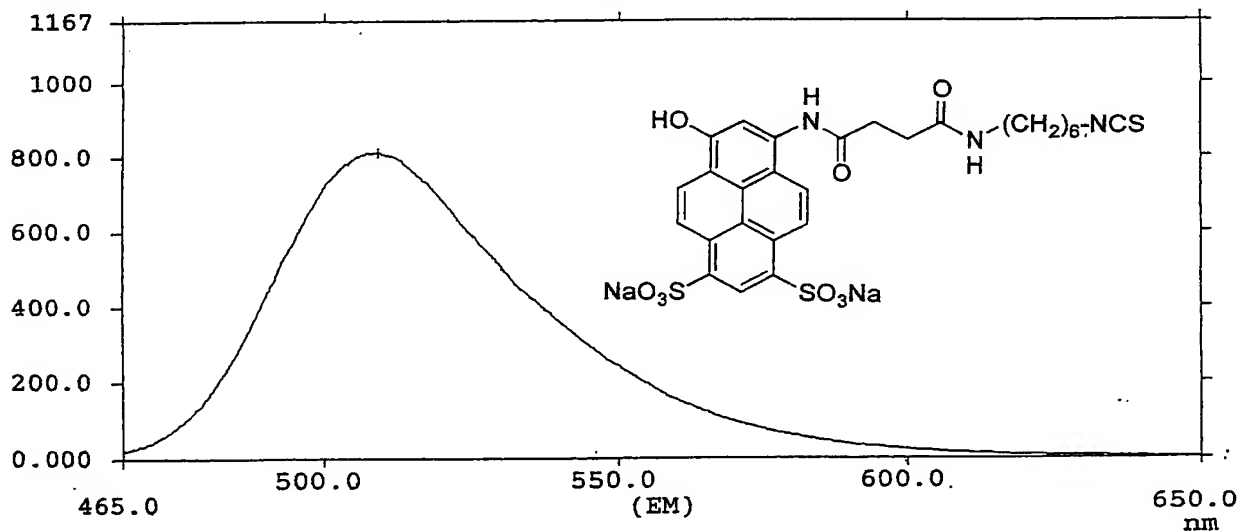
Fig. 1A



Sample : ABC-558-59
Comment : SB Susb.dil pH 9.0
EM : 508.0 nm
Data Mode : Fluorescence
Scan Speed : 1200 nm/min Slit (EX/EM) : 2.5 nm / 2.5 nm
PMT Voltage : 700 V Response : Auto

No.	WL (nm)	Peak	No.	WL (nm)	Peak
1	302.6	435.3	2	458.6	811.9

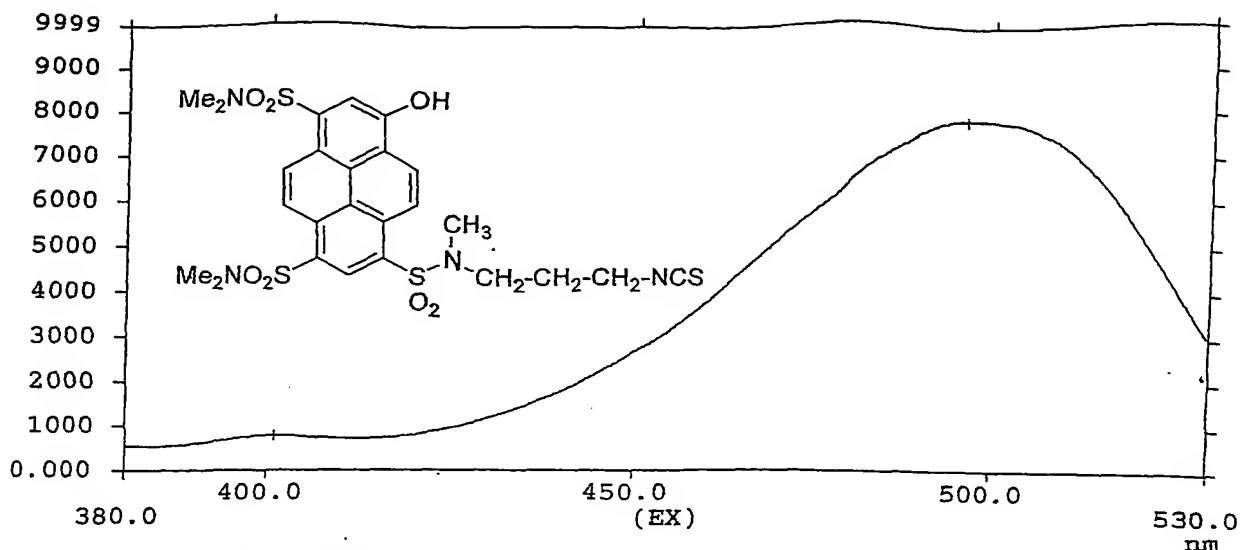
Data Peak Data 12/03/01 02:41 PM



Sample : ABC-558-59
Comment : SB Susb.dil pH 9.0
EX : 458.0 nm
Data Mode : Fluorescence
Scan Speed : 1200 nm/min Slit (EX/EM) : 2.5 nm / 2.5 nm
PMT Voltage : 700 V Response : Auto

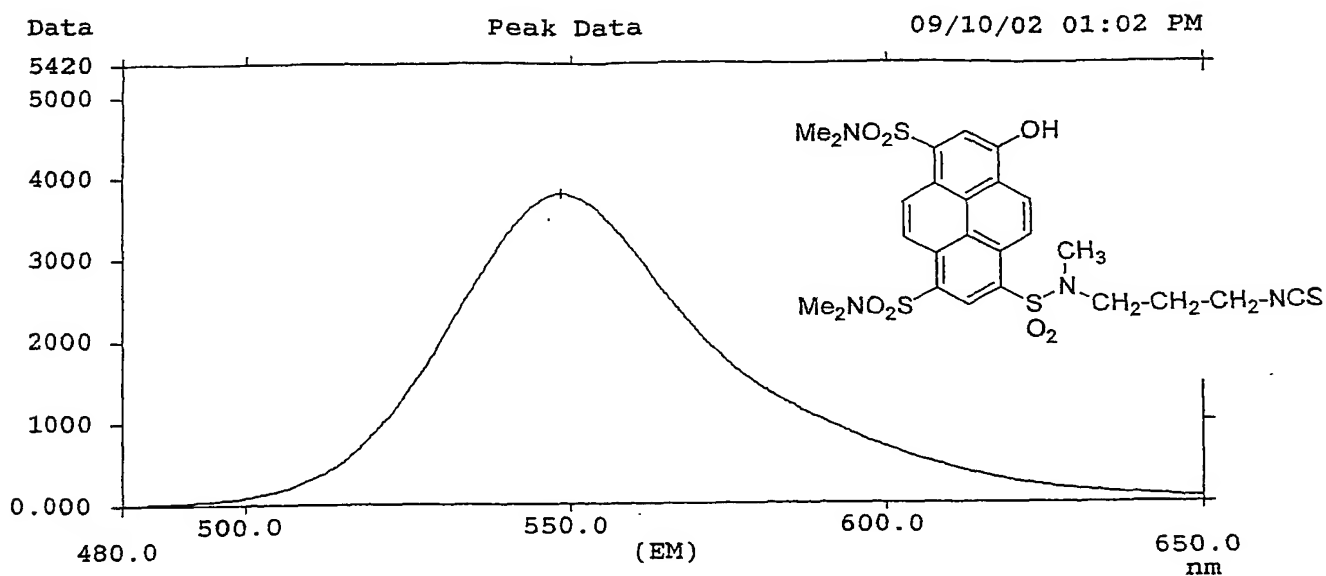
No.	WL (nm)	Peak	No.	WL (nm)	Peak
1	509.0	813.4			

Fig. 1B



Sample : SBO-R-NCS
 Comment : PBS pH 7.0
 EM : 547.0 nm
 Data Mode : Fluorescence
 Scan Speed : 1200 nm/min Slit (EX/EM) : 5.0 nm / 5.0 nm
 PMT Voltage : 700 V Response : Auto

No.	WL (nm)	Peak	No.	WL (nm)	Peak
1	401.0	783.5	2	496.2	7881



Sample : SBO-R-NCS
 Comment : PBS pH 7.0
 EX : 460.0 nm
 Data Mode : Fluorescence
 Scan Speed : 1200 nm/min Slit (EX/EM) : 5.0 nm / 5.0 nm
 PMT Voltage : 700 V Response : Auto

No.	WL (nm)	Peak	No.	WL (nm)	Peak
1	548.4	3807			

Fig. 2

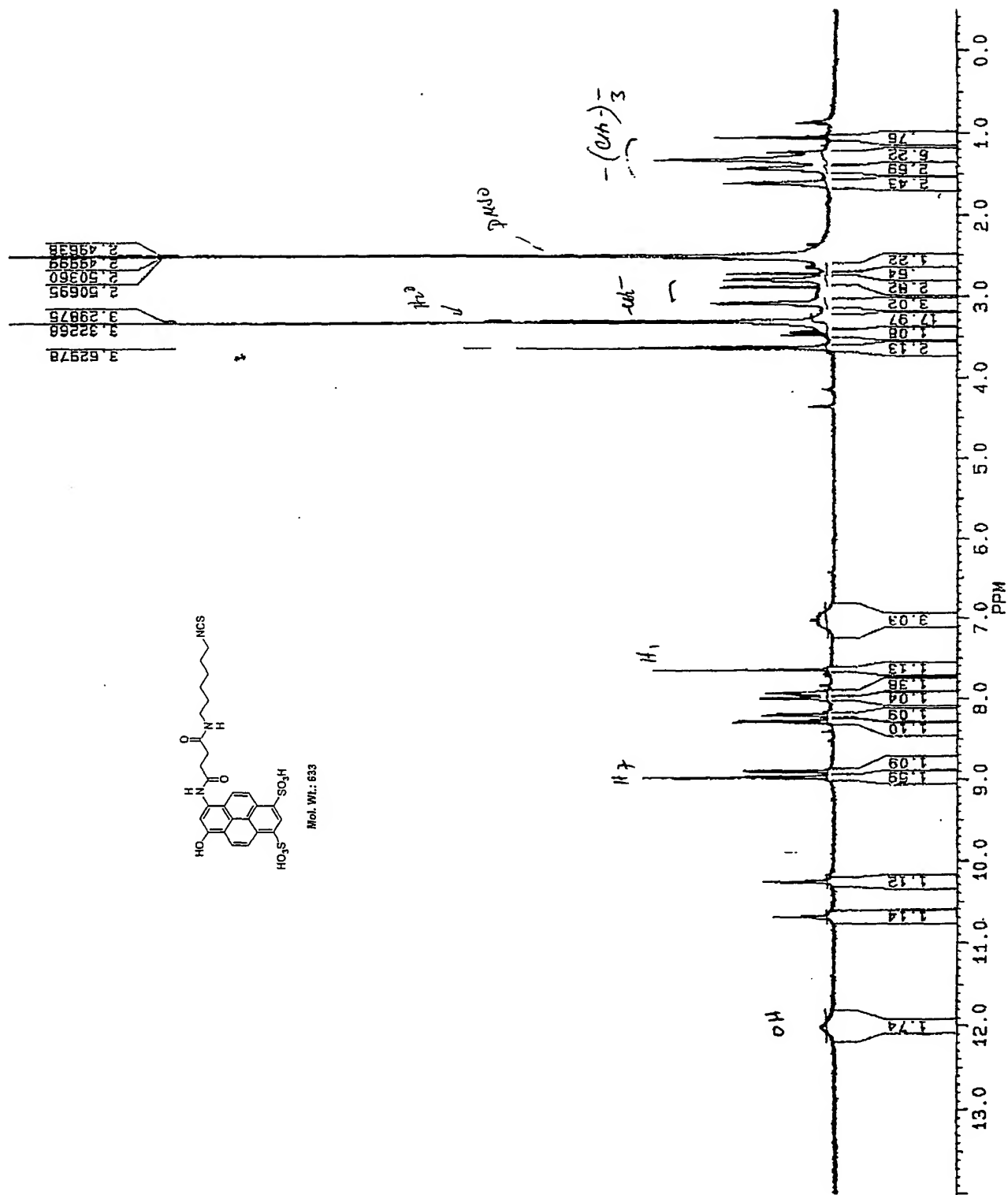


Fig. 3

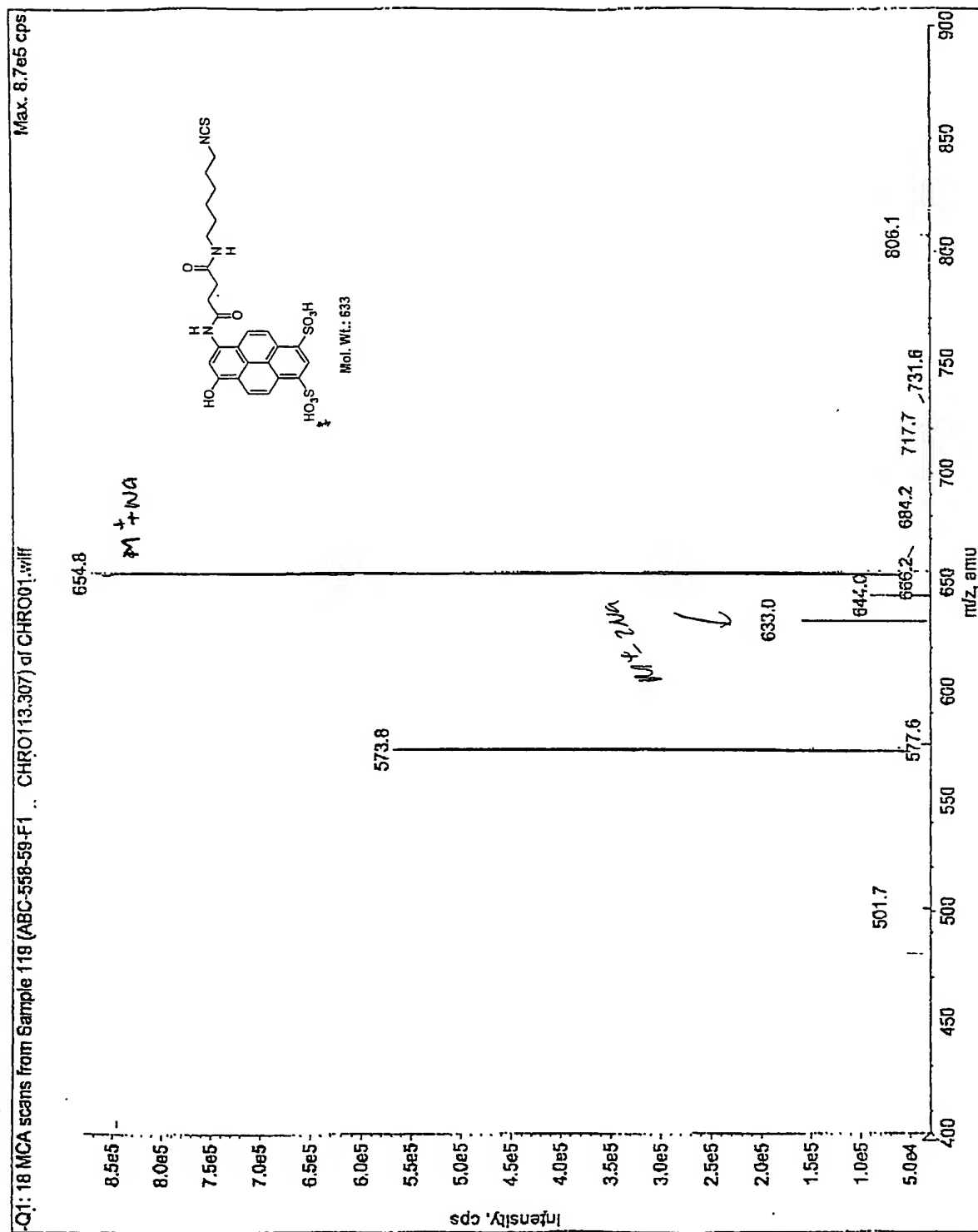


Fig. 4

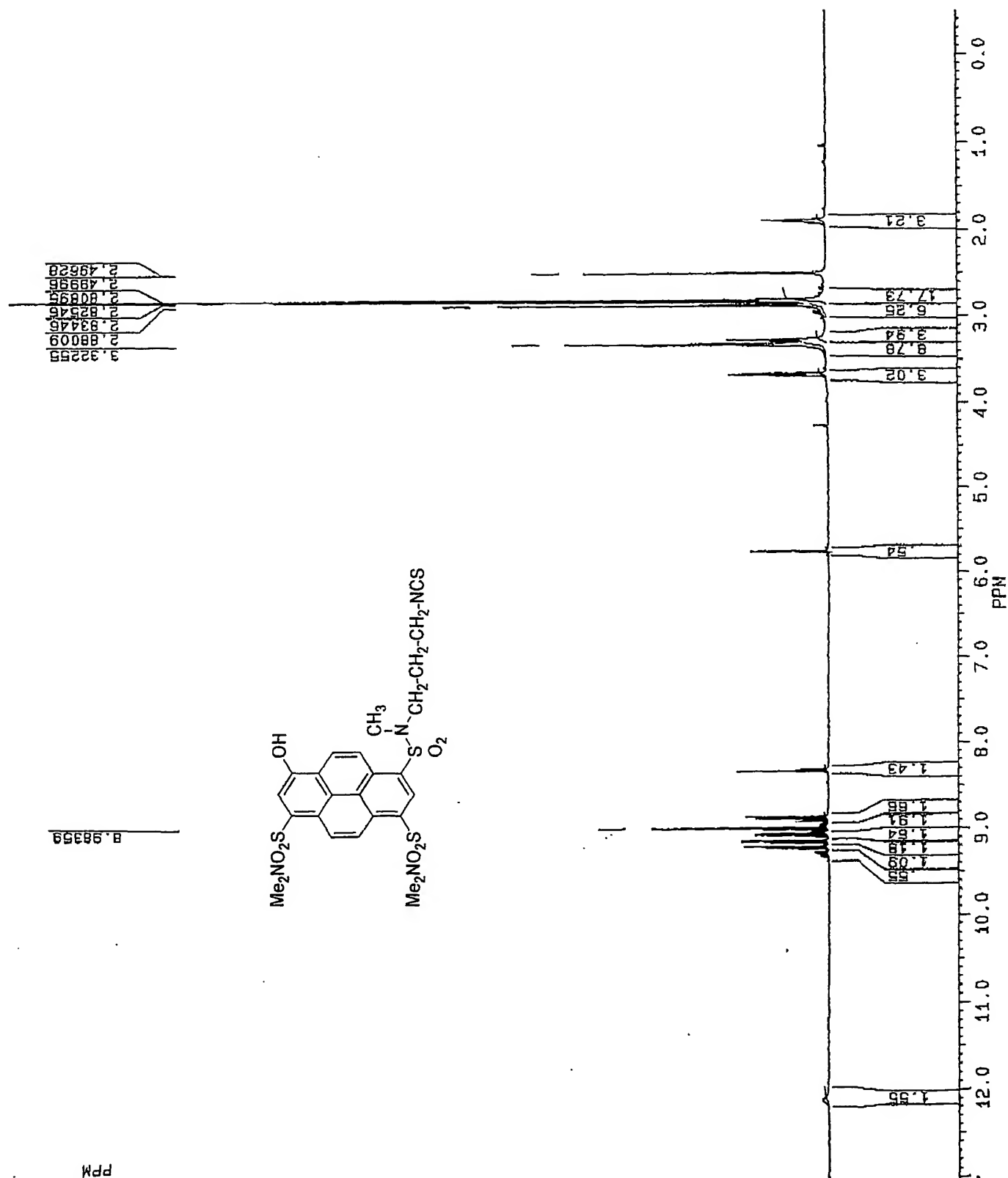
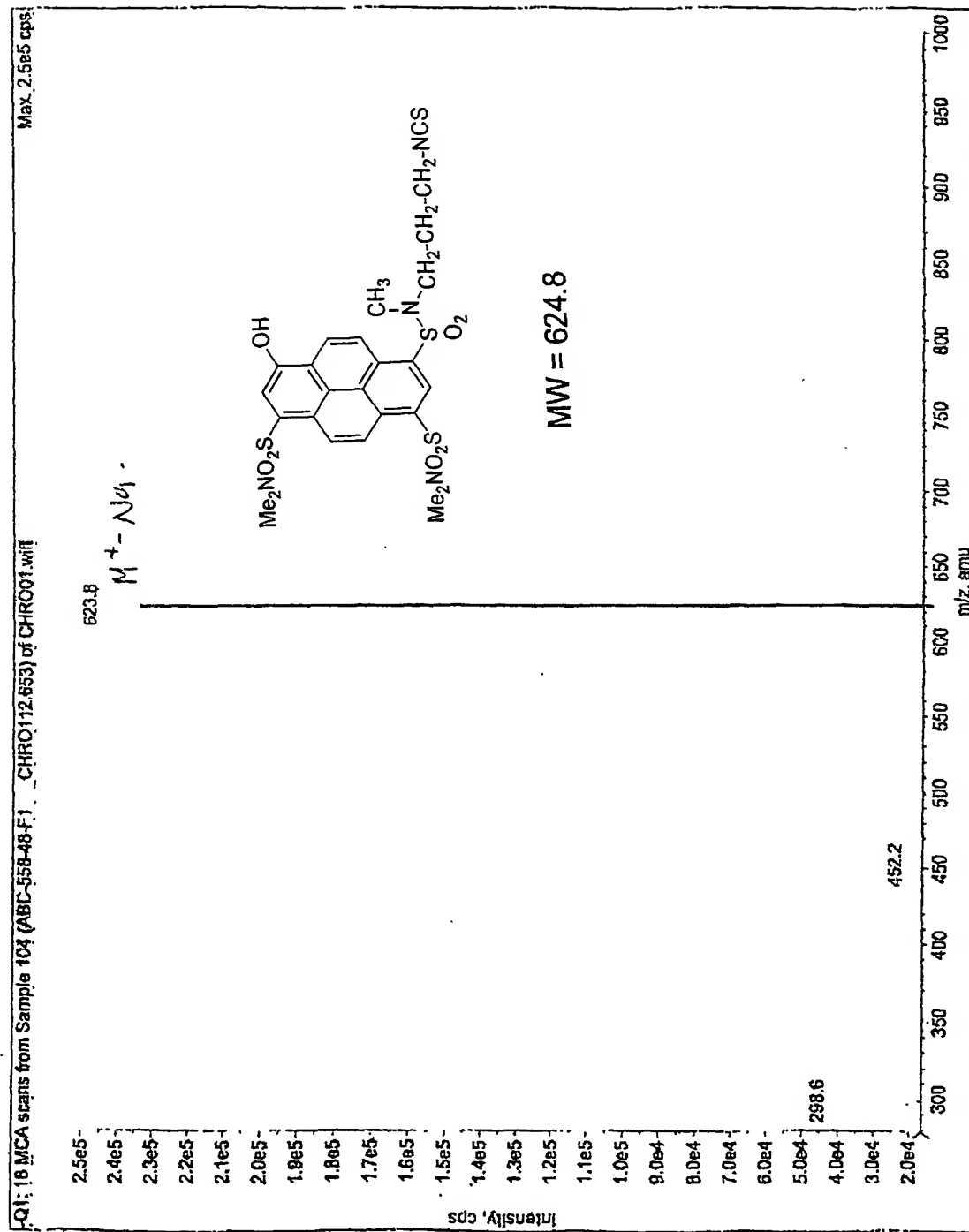


Fig. 5



NAME CHAN-LEV REP TYPE DIRECTORY
 COLLECTION DATA 59695F1 4 1 1 0119 C:\GOLD\ASCI\SAMPLE\ C:\GOLD\ASCI\METHOD\
 METHOD PEPT-REV
 SAMPLE TABLE 61901CPT C:\GOLD\CE811F\FAF54T7A3\ SYSTEM 1: SYSTEM1
 Chart Speed 0.50 cm/min

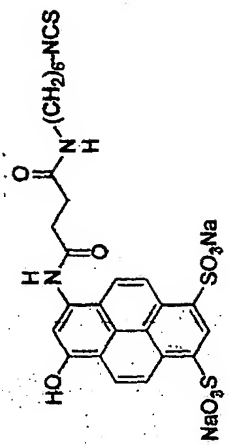
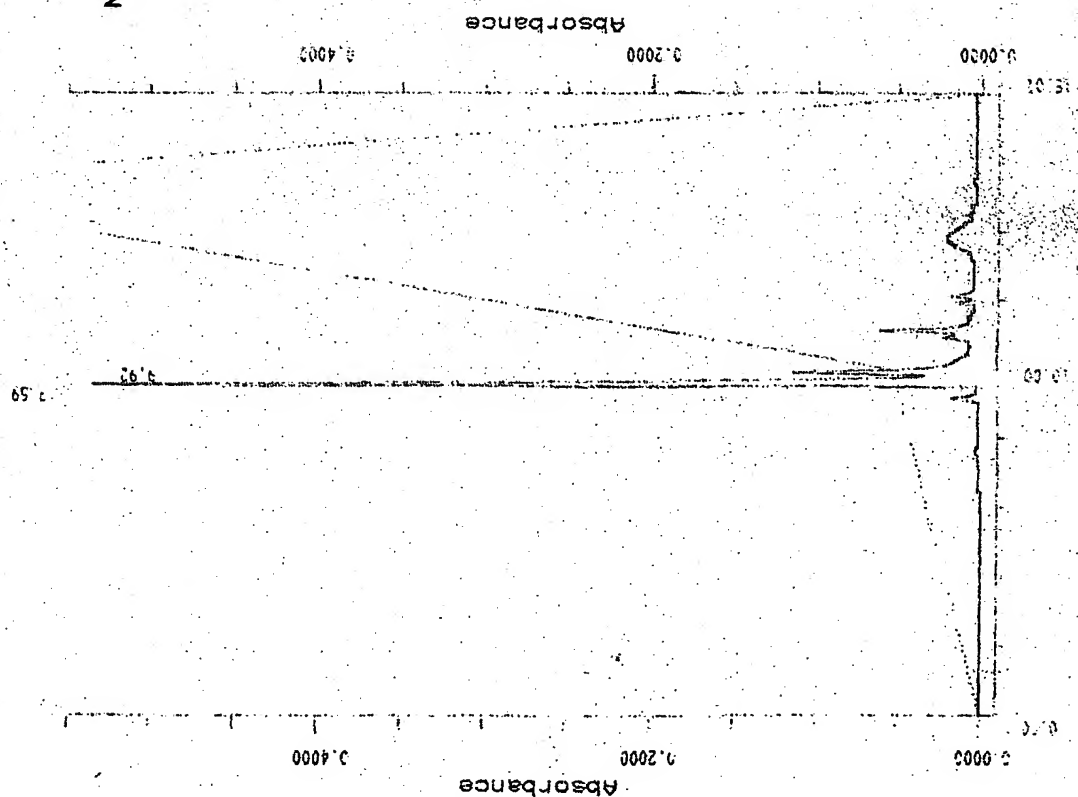
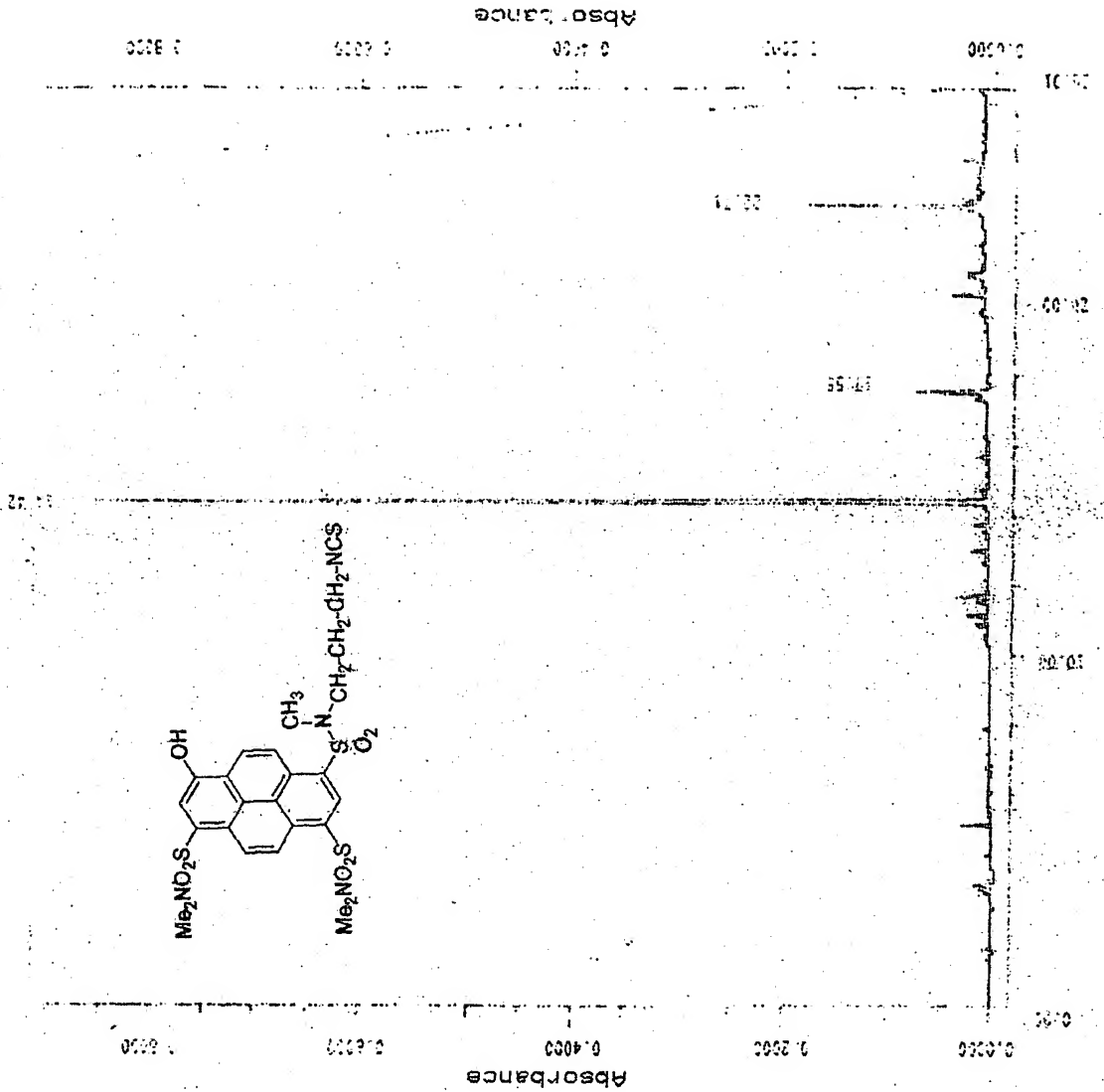


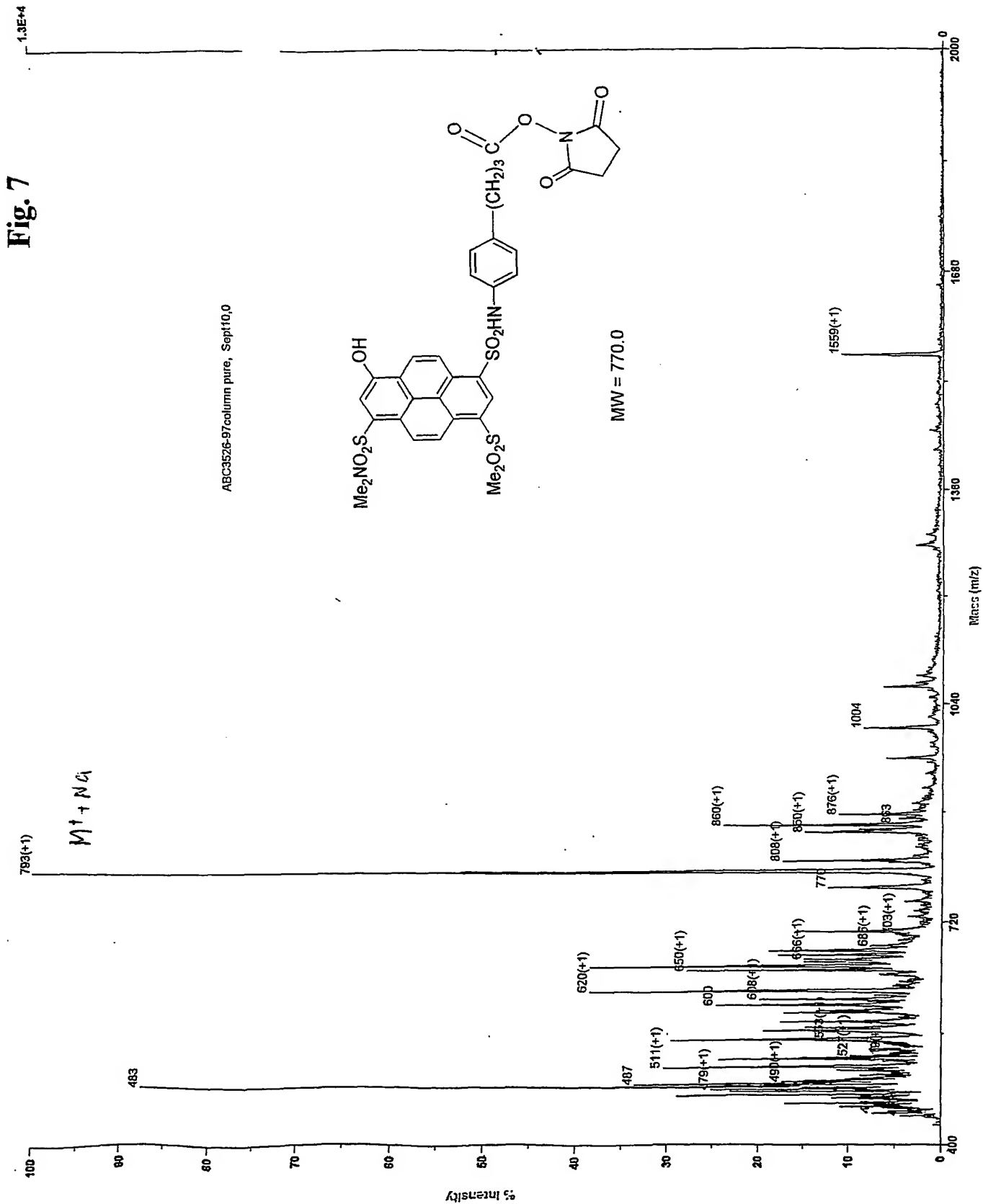
Fig. 6A

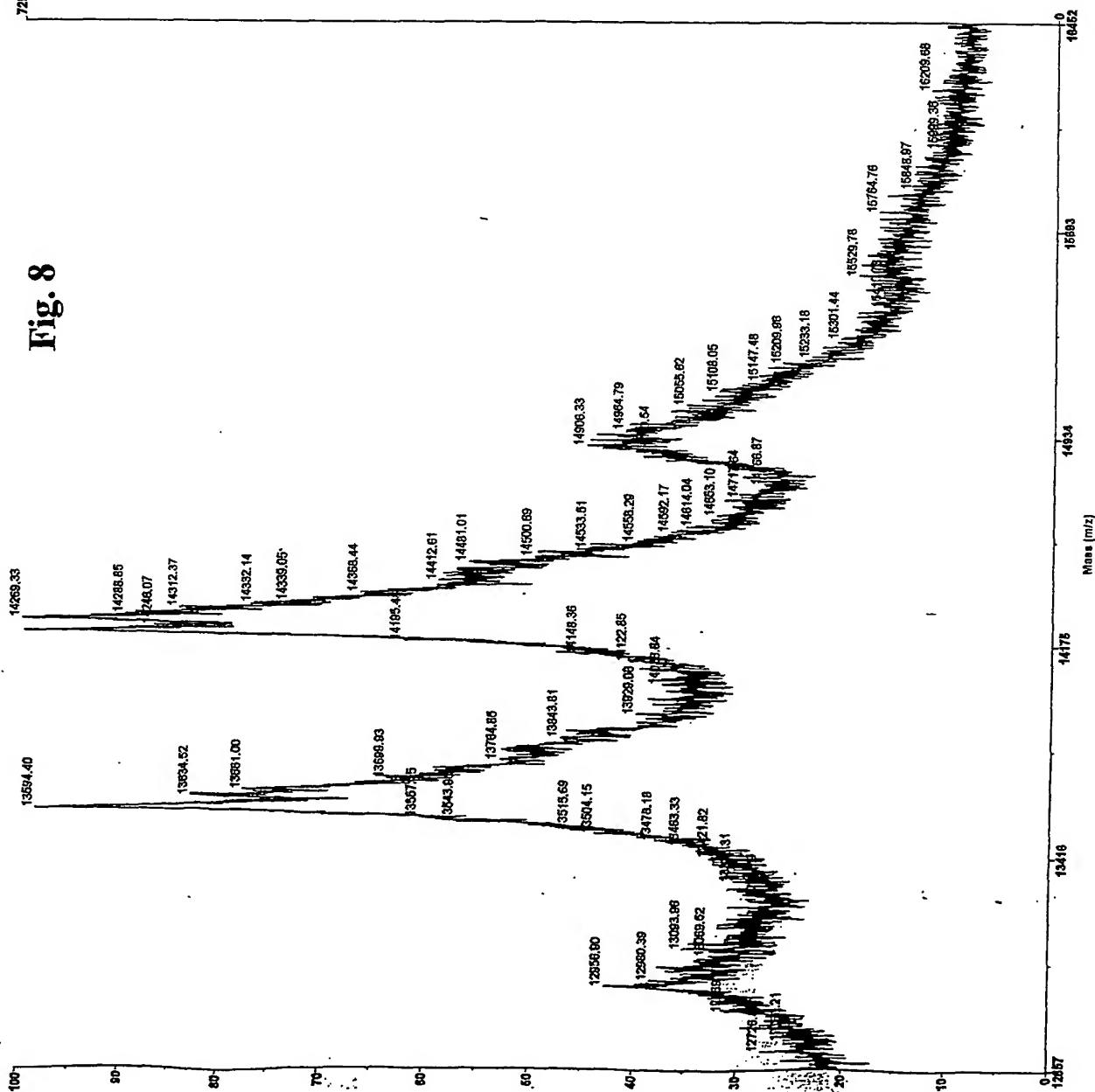
TIME DATE
 INJECTION 08:53:15 6-MAY 2002
 ANALYSIS 09:11:24 6-MAY 2002
 REPORT 09:11:26 6-MAY 2002

NAME CHAN LEV REP TYPE DIRECTORY
 COLLECTION DATA 9801C96 A 1 1 Orig C:\GOLD\ABC\SAMPLE\
 METHOD RAW'S C:\GOLD\ABC\METHOD\
 C:\GOLD\CEPTEK\OPIR126
 SYSTEM 1: SYSTEM1
 SAMPLE TABLE 61901CRT
 Chart Speed 0.50 cm/min
 Absorbance
 0.0000 0.2000 0.4000 0.6000 0.8000
 10.00 20.00 30.00 40.00 50.00 60.00 70.00 80.00 90.00 100.00
 INJECTION 10:18:12 7 AUG 2002
 ANALYSIS 10:44:22 7 AUG 2002
 REPORT 10:44:25 7 AUG 2002
 DATE TIME

Fig. 6B







Mode of operation: Linear
 Extraction mode: Delayed
 Polarity: Positive
 Acquisition control: Manual

Accelerating voltage: 25000 V
 Grid voltage: 93%
 Guide wire O: 0.15%
 Extraction delay time: 350 nsec

Acquisition mass range: 7500 - 25000 Da
 Number of laser shots: 50/spectrum
 Laser intensity: 2617
 Laser Rep Rate: 20.0 Hz
 Calibration type: Default
 Calibration matrix: Sinapinic acid
 Low mass gate: 7500 Da

Digitizer start time: 48.918
 Bin size: 2 nsec
 Number of data points: 20113
 Vertical scale: 200 mV
 Vertical offset: 0.2%
 Input bandwidth: 150 MHz

Sample well: 10
 Plate ID: 1
 Serial number: 1197
 Instrument name: Voyager-DE
 Plate type filename: C:\VOYAGER\100 well plate.plt
 Lab name: PE Biosystems

Absolute x-position: 47487.8
 Absolute y-position: 47081
 Relative x-position: 180.288
 Relative y-position: -226.471
 Shots in spectrum: 50
 Source pressure: 3.19e-007
 Mirror pressure: 0
 TC2 pressure: 0.01037
 TIS gate width: 30
 TIS flight length: 940

sbg-wash1 july25,02.bio - 1.000ml Overlaid Traces
 rbabcsbg-itc july 25,02.bio - 1.000ml
 rb644-39.bio - 1.000ml FRESH

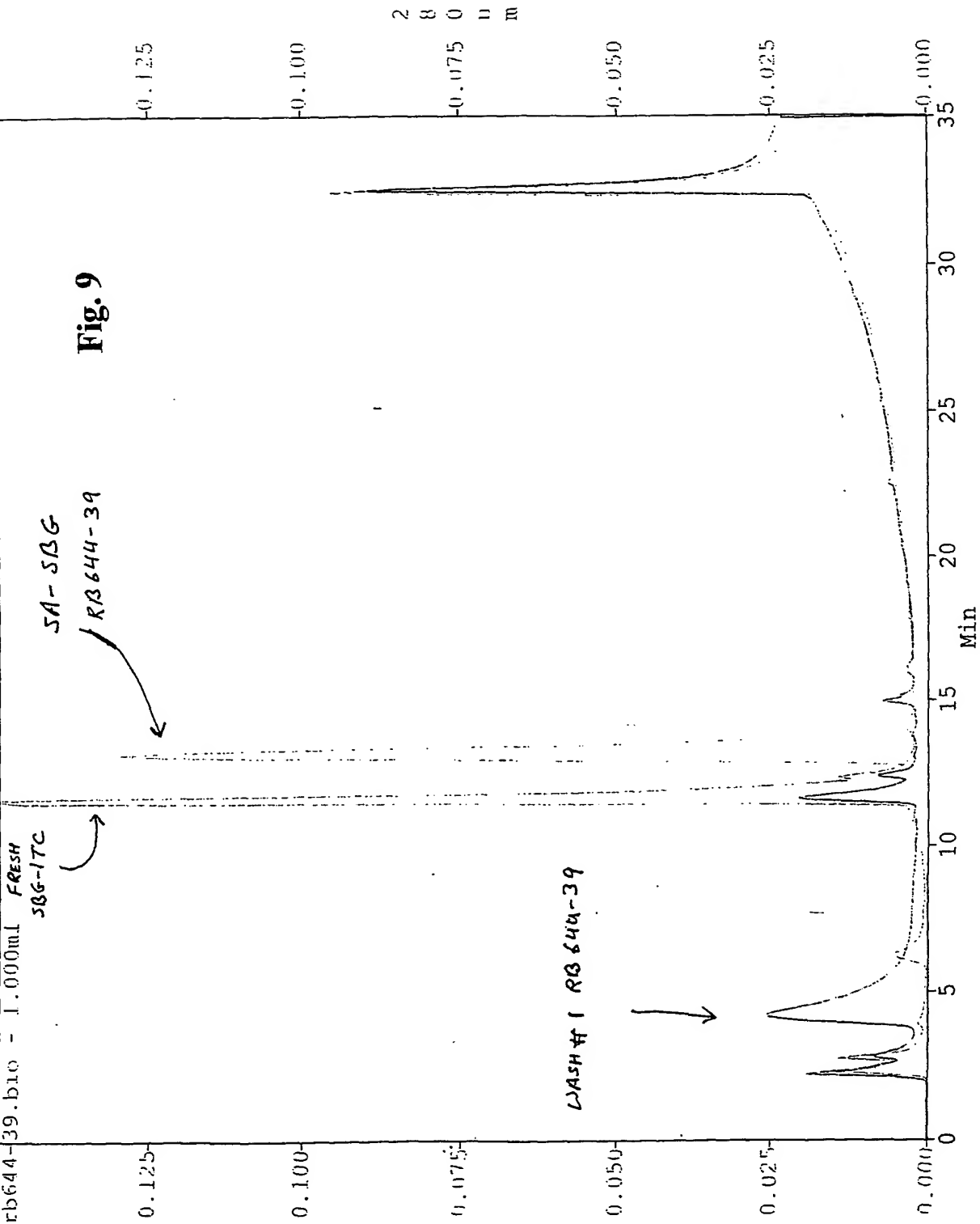
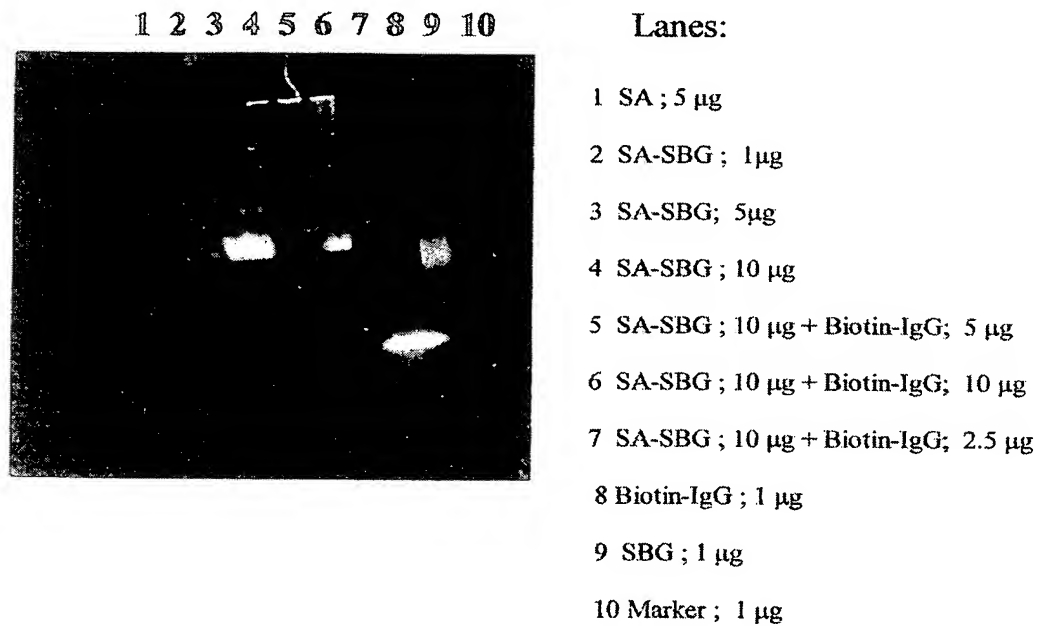


Fig. 9

Fig. 10

Gel Shift Assay of SA-SBG conjugate:



Mode of operation: Linear
 Extraction mode: Delayed
 Polarity: Positive
 Acquisition control: Manual

Accelerating voltage: 25000 V
 Grid voltage: 93%
 Guide wire 0: 0.2%
 Extraction delay time: 1700 nsec

Acquisition mass range: 10000 - 200000 Da
 Number of laser shots: 25/spectrum
 Laser intensity: 2817
 Laser Rep Rate: 20.0 Hz
 Calibration type: Default
 Calibration matrix: Sinapinic acid
 Low mass gate: 10000 Da

Digitizer start time: 56.65
 Bin size: 10 nsec
 Number of data points: 19515
 Vertical scale: 200 mV
 Vertical offset: 0.5%
 Input bandwidth: 150 MHz

Sample well: 32
 Plate ID: 1
 Serial number: 1197
 Instrument name: Voyager-DE
 Plate type filename: C:\VOYAGER\100 well plate.plt
 Lab name: PE Biosystems

Absolute x-position: 7283.04
 Absolute y-position: 31932.2
 Relative x-position: 615.544
 Relative y-position: -135.304
 Shots in spectrum: 25
 Source pressure: 3.723e-007
 Mirror pressure: 0
 TC2 pressure: 0.00989
 TIS gate width: 30
 TIS flight length: 940

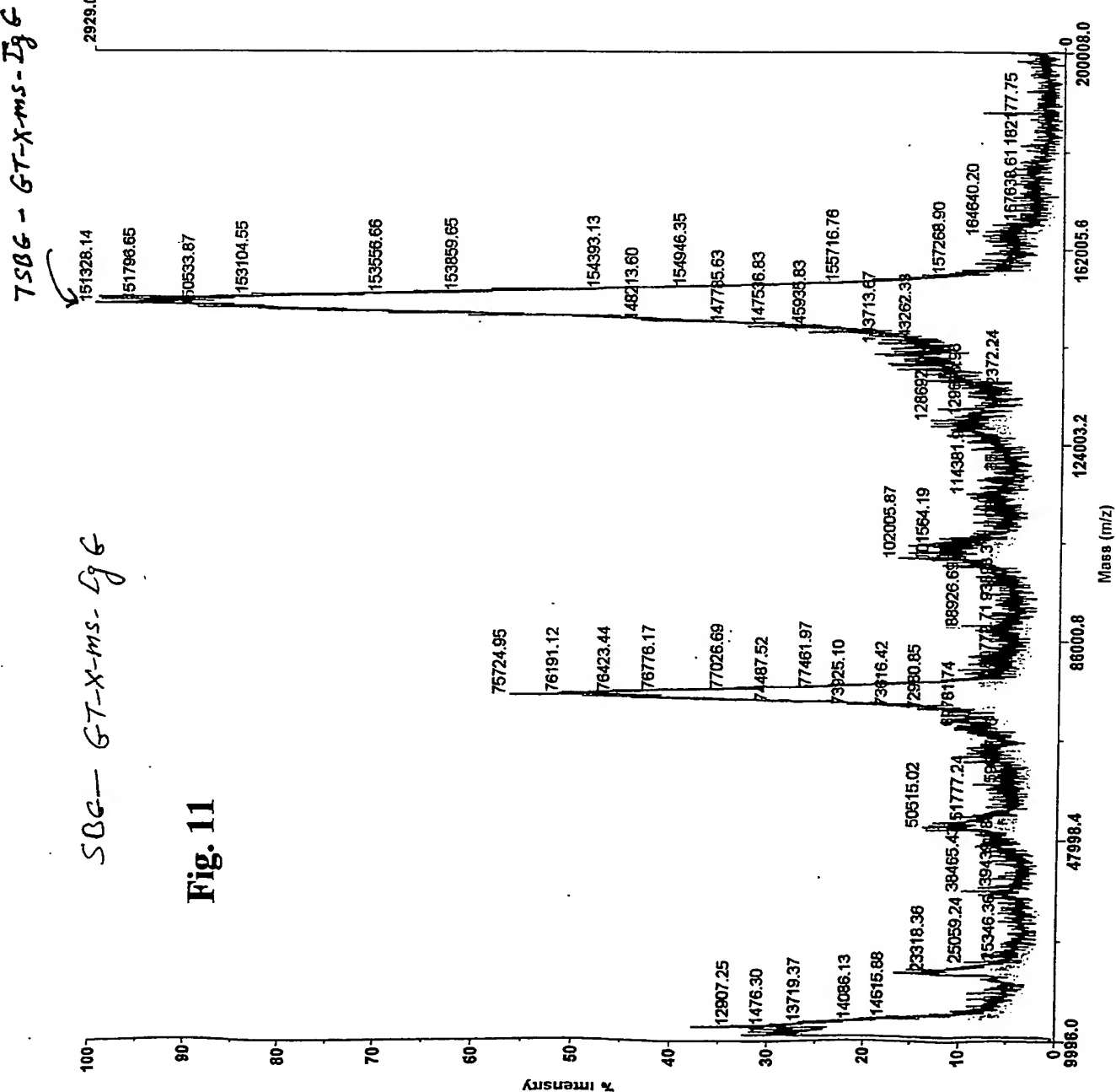
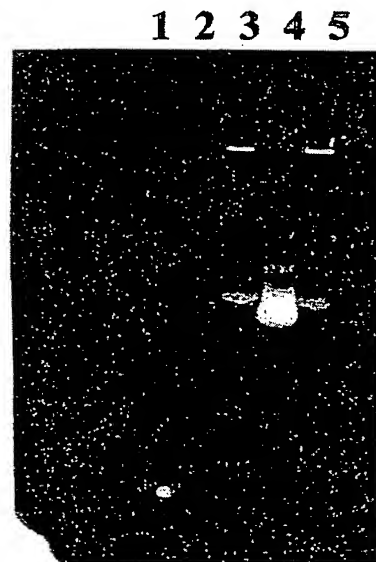


Fig. 12

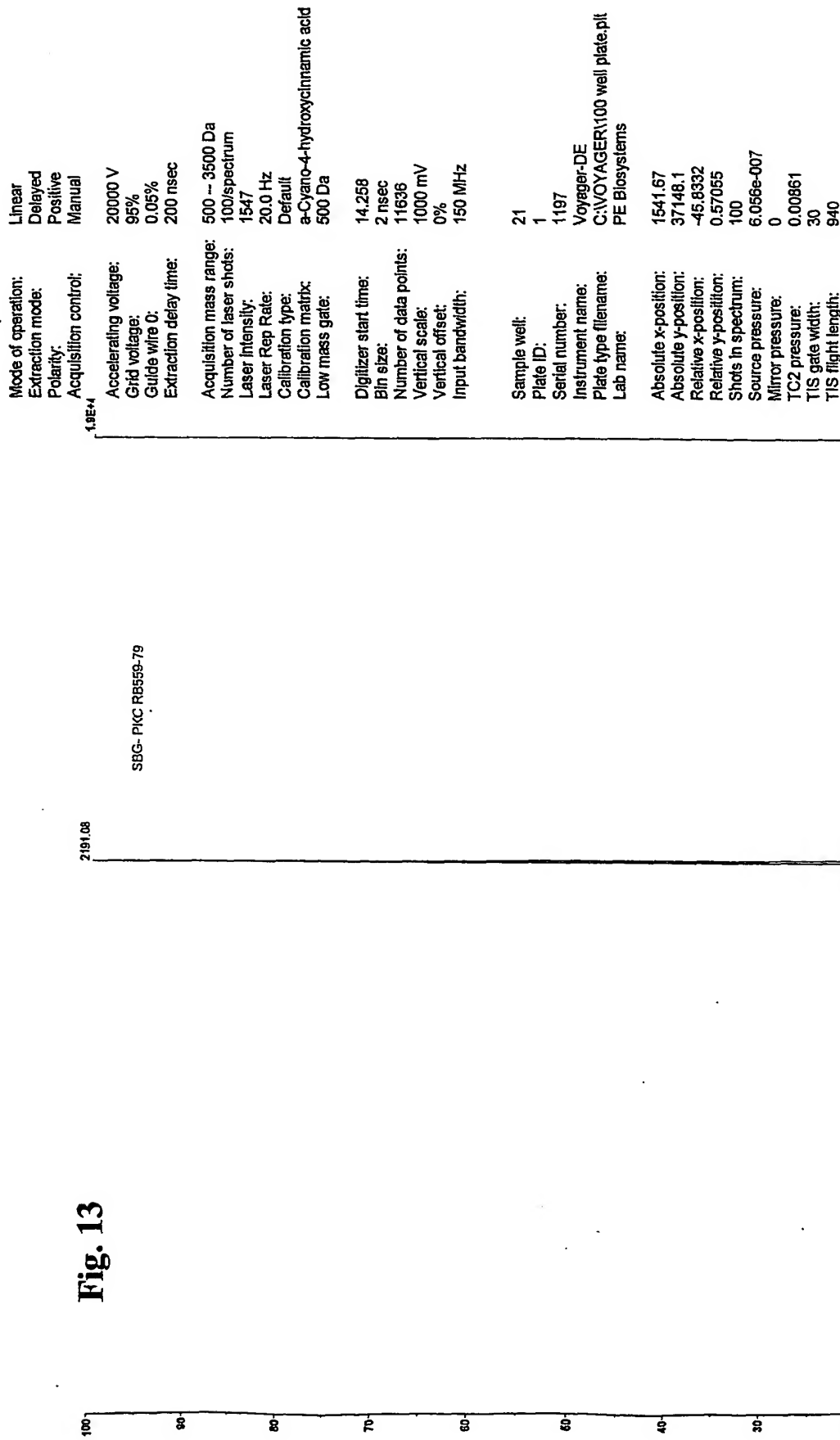
Gel Shift Assay of SA-SBO



Lanes:

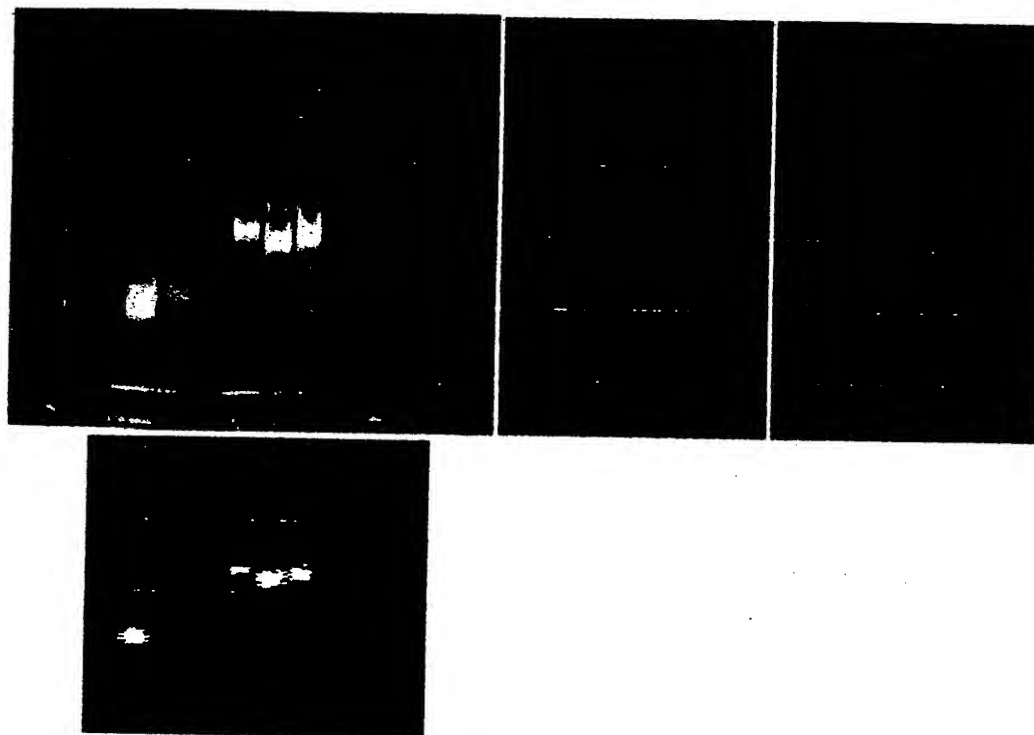
1. SA-SBO ; 1 μ g
2. SA-SBO + Biotin-IgG ; 5 μ g
3. SA-SBO ; 10 μ g
4. SA-SBO + Biotin-IgG ; 10 μ g
5. Biotin-IgG ; 5 μ g

Fig. 13



POLAROID PHOTOGRAPH

**DIGITAL IMAGES
at two thresholds**



**Digital Image showing
Lane alignment**

Figure 14 is a digital image of a polyacrylamide gel showing fluorescent conjugates formed by labeling streptavidin and IgG molecules with the isothiocyanate of StarBright Orange to give labeled reporter moieties having measurable label to probe ratios.

Fig. 15



Figure 15 is a photograph of a polyacrylamide gel showing the fluorescence of an oligonucleotide labeled with StarBright Green Dye.

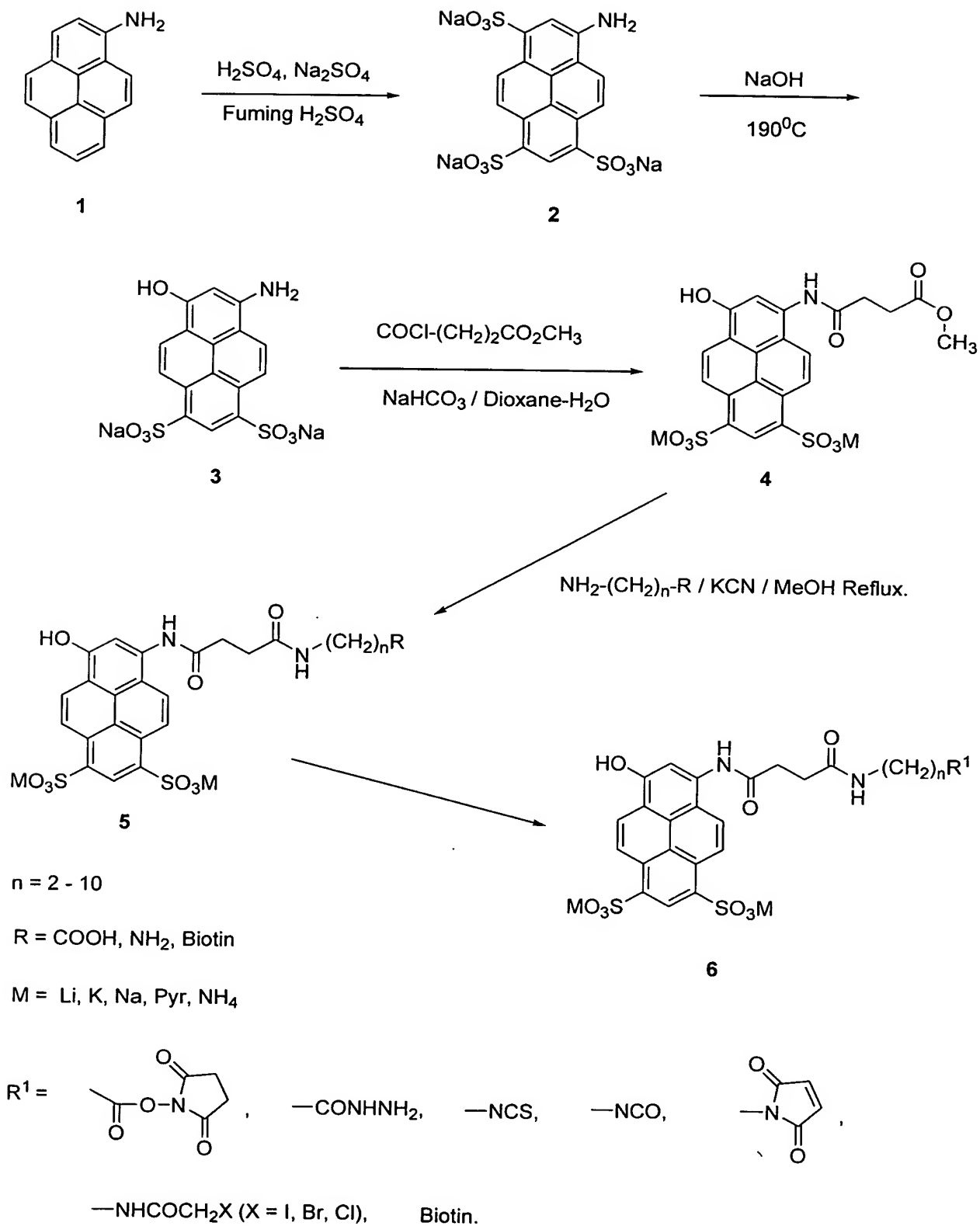


FIG. 16

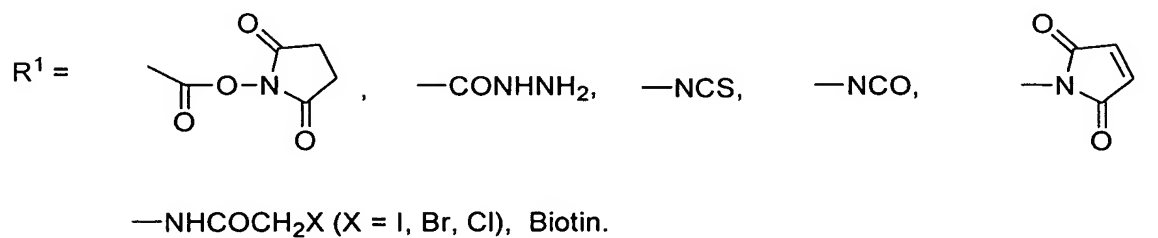
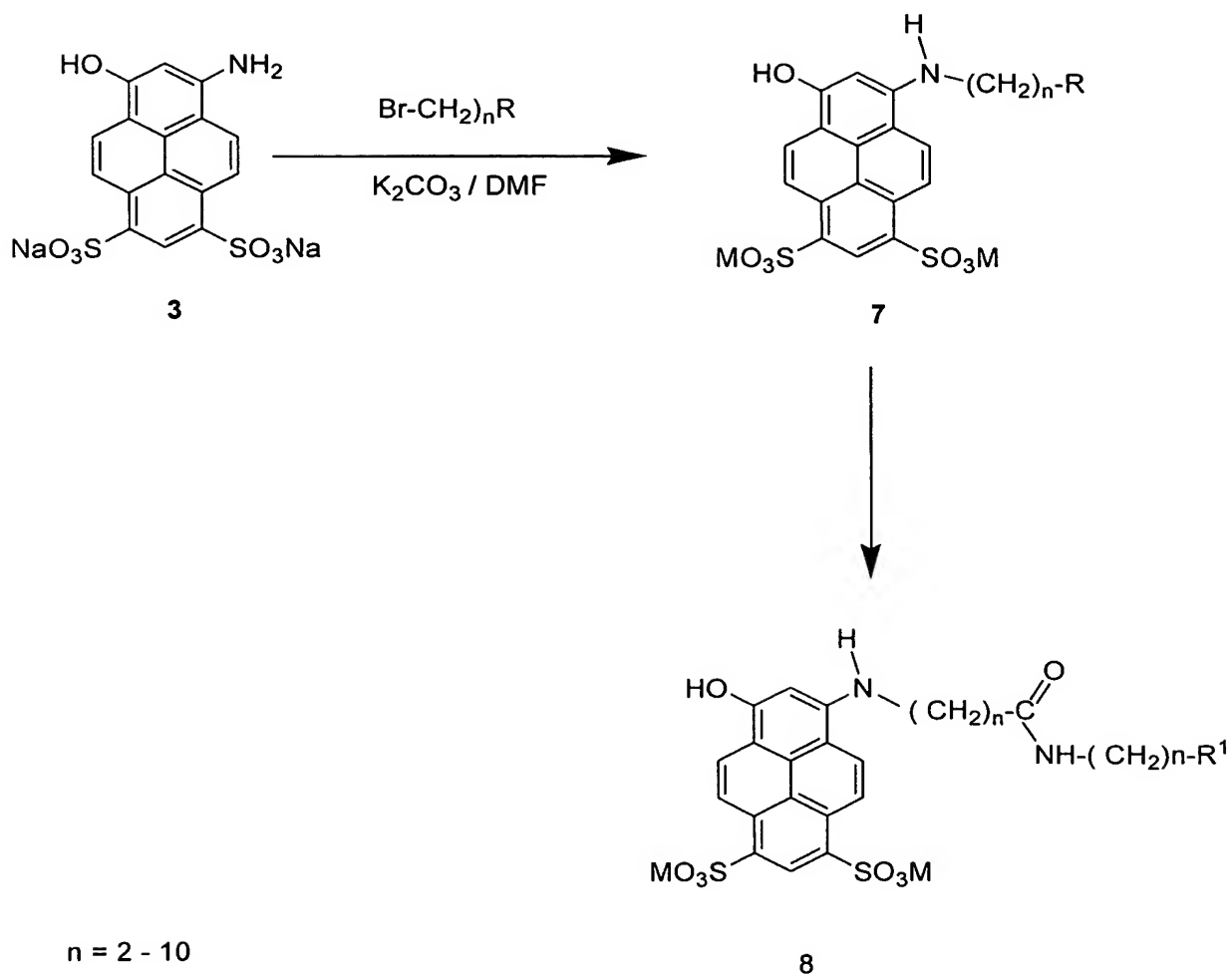


FIG. 17

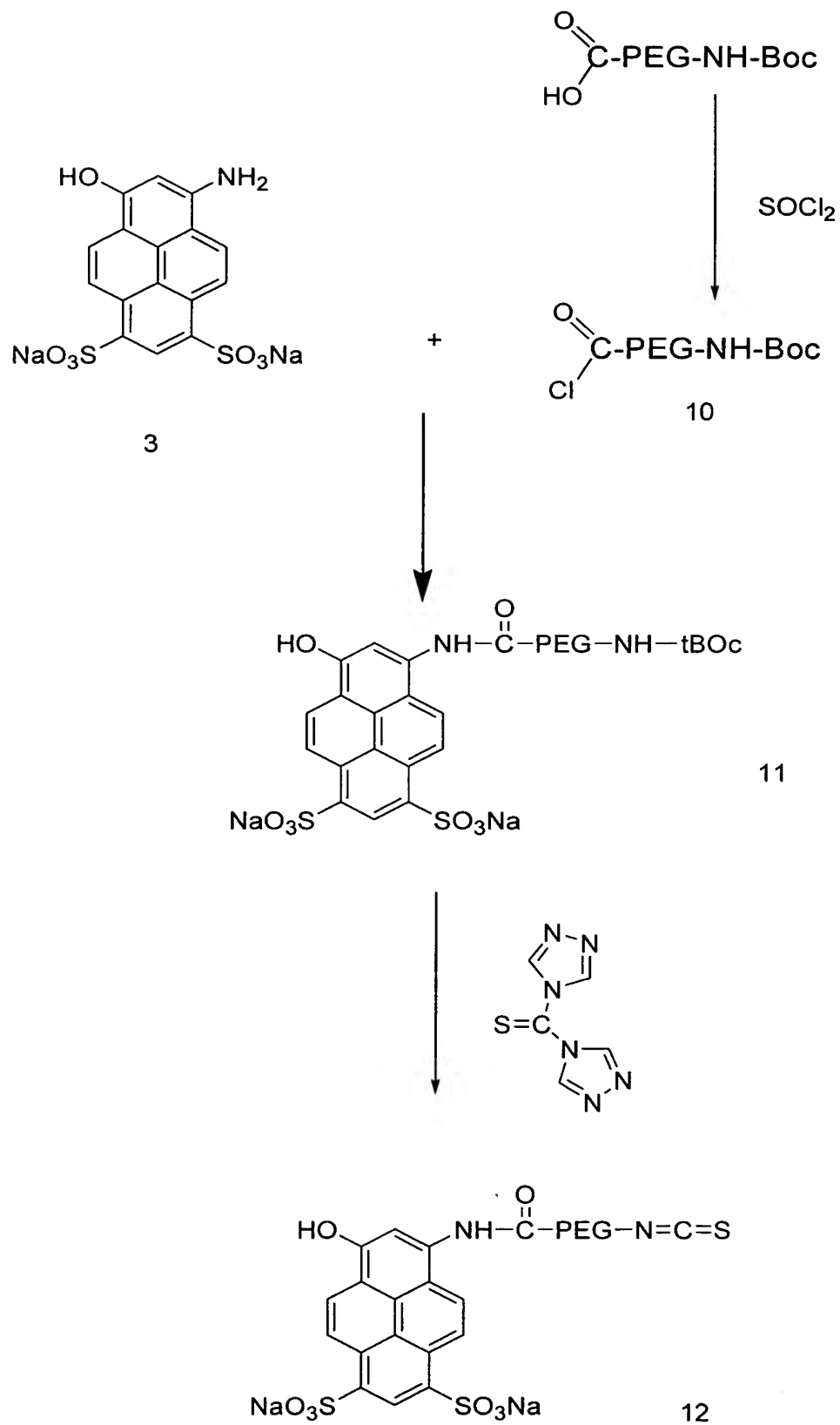
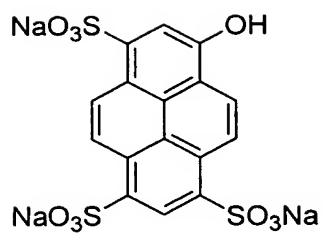
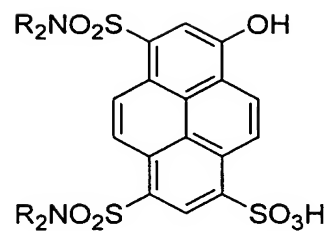
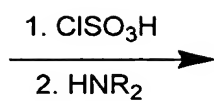


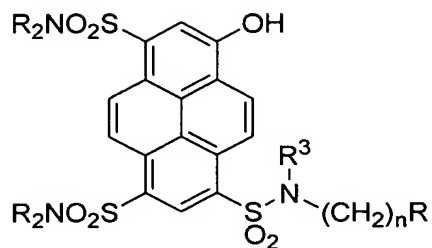
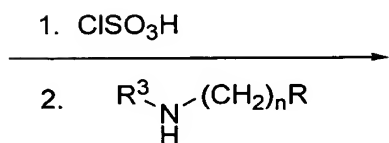
FIG. 18



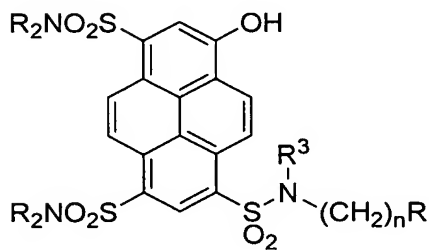
13



14



15



16

$n = 2 - 10$

$\text{R}^1, \text{R}^2 = \text{alkyl groups}$

$\text{R}^3 = \text{H, alkyl groups, } \text{R}^4 = \text{COOH, NH}_2, \text{Biotin}$

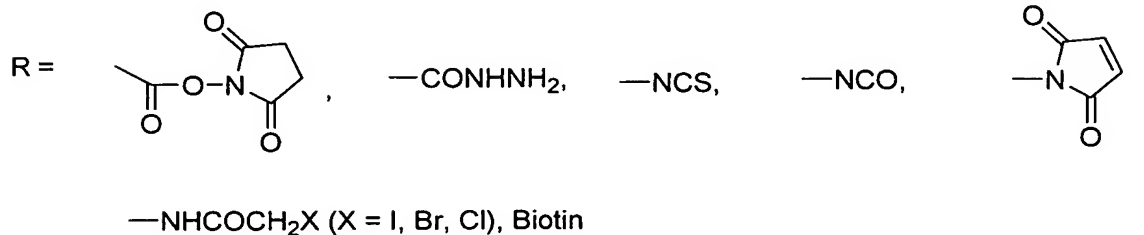
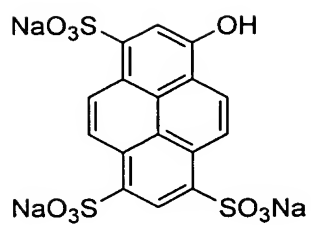
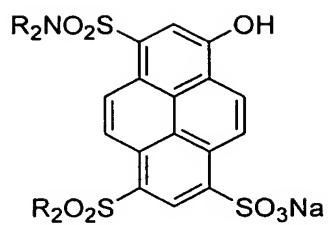
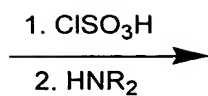


FIG. 19



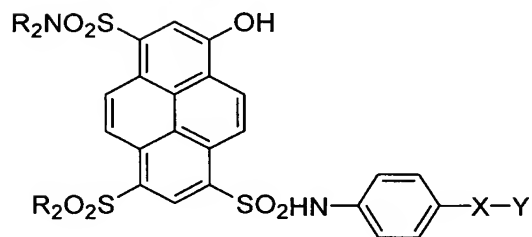
13



14

1. ClSO_3H

2. $\text{H}_2\text{N}-\text{C}_6\text{H}_4-\text{X}-\text{Y}$
Pyridine

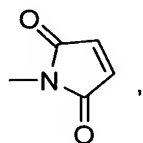
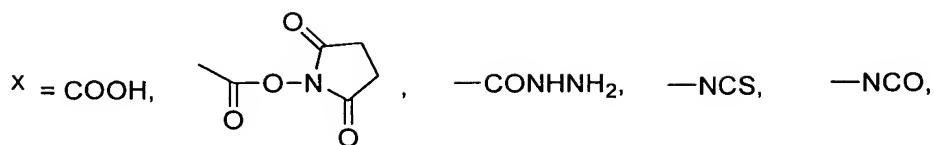


17

$n = 0 - 8$

R = Alkyl groups

$\text{X} = -(\text{CH}_2)_n-$



$-\text{NHCOCH}_2\text{X}$ ($\text{X} = \text{I}, \text{Br}, \text{Cl}$), Biotin

FIG. 20

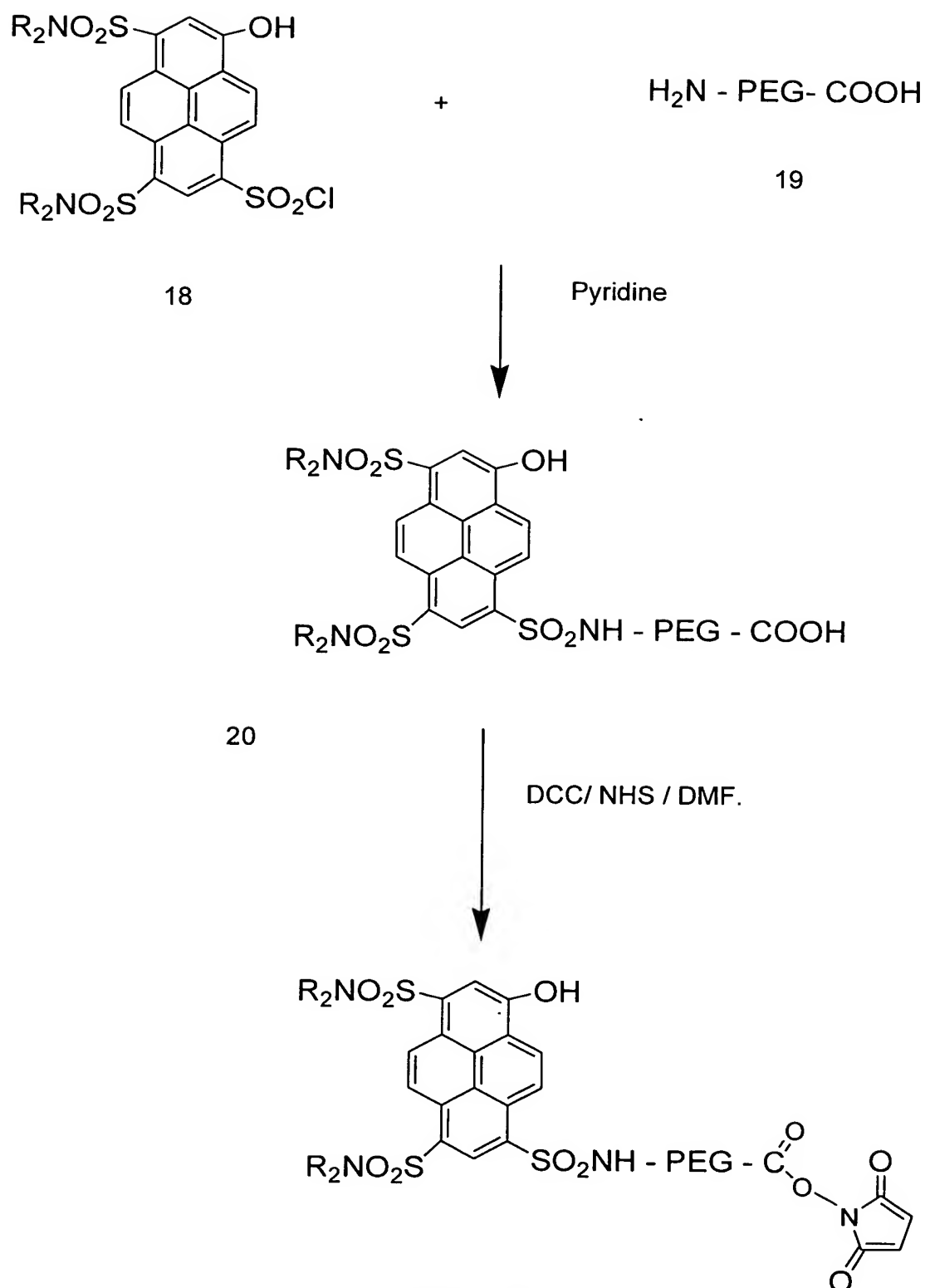


FIG. 21